

THE PASSAGE OF THE BOMB RADIOCARBON PULSE INTO THE PACIFIC OCEAN

William J Jenkins¹ • Kathryn L Elder • Ann P McNichol • Karl von Reden

National Ocean Sciences Accelerator Mass Spectrometer Facility, Woods Hole Oceanographic Institution, Woods Hole, Massachusetts 02543, USA.

ABSTRACT. We report and compare radiocarbon observations made on 2 meridional oceanographic sections along 150°W in the South Pacific in 1991 and 2005. The distributions reflect the progressive penetration of nuclear weapons-produced ¹⁴C into the oceanic thermocline. The changes over the 14 yr between occupations are demonstrably large relative to any possible drift in our analytical standardization. The computed difference field based on the gridded data in the upper 1600 m of the section exhibits a significant decrease over time (approaching 40 to 50‰ in $\Delta^{14}\text{C}$) in the upper 200–300 m, consistent with the decadal post-bomb decline in atmospheric ¹⁴C levels. A strong positive anomaly (increase with time), centered on the low salinity core of the Antarctic Intermediate Water (AAIW), approaches 50–60‰ in $\Delta^{14}\text{C}$, a clear signature of the downstream evolution of the ¹⁴C transient in this water mass. We use this observation to estimate the transit time of AAIW from its “source region” in the southeast South Pacific and to compute the effective reservoir age of this water mass. The 2 sections show small but significant changes in the abyssal ¹⁴C distributions. Between 1991 and 2005, $\Delta^{14}\text{C}$ has increased by 9‰ below 2000 m north of 55°S. This change is accompanied overall by a modest increase in salinity and dissolved oxygen, as well as a slight decrease in dissolved silica. Such changes are indicative of greater ventilation. Calculation of “phosphate star” also indicates that this may be due to a shift from the Southern Ocean toward North Atlantic Deep Water as the ventilation source of the abyssal South Pacific.

INTRODUCTION

Observation of the passage of anthropogenically produced transient tracers into the oceans has provided us with an opportunity to characterize ocean ventilation, circulation, and mixing in new ways. In the simplest sense, the penetration of recently produced tracers such as CFCs or anthropogenic radioisotopes into the ocean basins creates a qualitative visualization of the pathways (e.g. Jenkins and Rhines 1980) and timescales (e.g. Schlosser et al. 2001) associated with water mass formation and fluid motion. As importantly, these observations can be effectively used to diagnose the performance of large-scale ocean models (England and Maier-Reimer 2001; Doney and Hecht 2002) or in inversions aimed at the estimation of oceanic inventories of anthropogenic carbon (McNeil et al. 2003; Waugh et al. 2006). Such tracers offer a range of characteristics including how they enter the ocean, their biogeochemical behavior in the marine environment, their half-lives, and their time history at the sea surface. Consequently, each tracer tells us something different about ocean ventilation and circulation.

Radiocarbon is an important tracer, especially in the Pacific, because the distribution of natural ¹⁴C offers information on the longest (century to millennium) timescales in the deep waters (Munk 1966; Craig 1969), while the bomb-produced transient component informs us of processes operating on decadal timescales (Broecker and Peng 1980; Mahadevan 2001; Rodgers et al. 2004). Where the bomb and natural ¹⁴C distributions overlap, the story becomes more complex, and strategies for separating these 2 components have been proposed (Broecker et al. 1995; Rubin and Key 2002). Perhaps the most powerful diagnostic, however, is the observation of the temporal evolution of ¹⁴C distribution; to a first approximation, the natural component will not vary while the bomb transient necessarily evolves.

The GEOSECS expeditions of the 1970s offered the first large-scale glimpse of the ¹⁴C distribution in the Pacific (Bainbridge et al. 1987; Ostlund et al. 1974), which was subsequently mapped in

¹Corresponding author. Email: wjenkins@whoi.edu.

greater detail by the World Ocean Circulation Experiment (WOCE) large-scale survey in the 1990s (Key 1996; Key et al. 1996, 2002). After this, the CLIVAR (Climate Variability and Predictability) program has organized a series of repeat section occupations at approximately decadal intervals to monitor climate-induced changes in the ocean and to document the evolution of a number of transient tracers in the water column.

We report here a comparison of 2 ^{14}C sections taken along 150°W in the South Pacific between approximately 20°S and 63°S ; one taken in 1991 during the WOCE global survey and one occupied in 2005 as part of the CLIVAR Repeat Hydrography Program. In the next section, we provide a brief description of our methods along with quality control procedures used to establish the fidelity of our results, particularly on decadal timescales. After that, we describe and interpret the observed changes in shallow and intermediate waters. Finally, we show changes in the deep water and attempt to link those to concomitant changes in hydrography.

METHODS

Samples were collected using standard WOCE protocols whereby water was transferred from rosette-mounted Niskin bottles to 0.5-L borosilicate bottles with ground glass stoppers and “fixed” with saturated mercuric chloride solution to arrest biological activity (Key 1996). On return to the laboratory, the dissolved inorganic carbon (DIC) was removed in a vacuum system by acidification, sparging with ultrapure nitrogen, cryogenically purified, and stored as CO_2 in glass ampoules (McNichol et al. 1994). A portion of this gas was graphitized using hydrogen reduction over iron (Vogel et al. 1987; McNichol et al. 1992), and the graphite pressed into aluminum targets and analyzed by cesium sputtering accelerator mass spectrometry (AMS). The unused sample gas was archived for possible future analysis.

The AMS ^{14}C analyses were standardized against the NIST oxalic acid standards (OXI and OXII), and measurement performance was monitored by regular determinations of secondary (“consensus”) standards of varying ^{14}C activity. Since its inception, the laboratory has participated in all of the international intercalibration exercises, the most recent of which is VIRI (Scott et al. 2003), where our intercalibration results consistently fall within the range circumscribed by the major AMS measurement facilities.

Results here are reported in the standard $\Delta^{14}\text{C}$ format (Stuiver and Polach 1977), and the experimental uncertainty, estimated from ^{14}C ion counting statistics, comparison of replicate seawater samples, and reproducibility of primary and secondary standards, is better than 5‰ in $\Delta^{14}\text{C}$ on individual samples (Key 1996; Key et al. 1996; Elder et al. 1998).

One challenge when interpreting time-series measurements is to rule out potential drift in analytical standardization as a contributor to observed trends in the data. As an additional precaution, we later remeasured archived gas samples from WOCE stations to check for this. This approach will quantify any changes due to graphitization and AMS analysis. Figure 1 is a plot of just such a comparison for WOCE station P16S 198 ($28^\circ29.8'\text{S}$, $150^\circ29.9'\text{W}$, 19 August 1991) for the same samples measured in 1994 and again in 2007. The average offset (2007–1994) is $-0.95 \pm 1.13\text{‰}$, with a standard deviation of 5.75‰ . The latter quantity is consistent with our published uncertainties, as its expectation value is $\sqrt{2} \sigma$. As importantly, the remeasurement of the 1991 WOCE samples occurred in the same timeframe as our measurement of the samples from the 2005 occupation of P16S, and identical laboratory and analytical procedures were employed with both sets of samples. In any case, the mean offset obtained from this comparison provides us with a firm upper bound of 2‰ on any systematic effects that may contribute to any observed differences between the 2 cruise occupations.

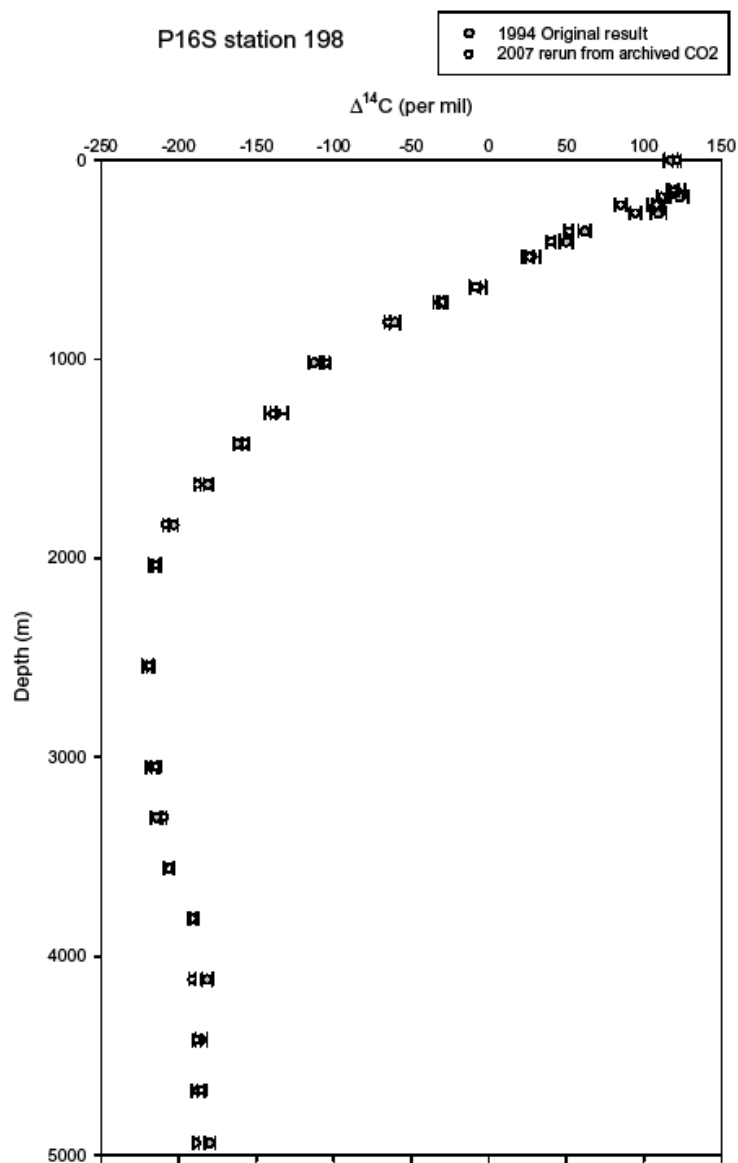


Figure 1 A comparison of WOCE DI^{14}C measured on a South Pacific WOCE station taken in 1991 measured at 2 points in time: 1994 (filled circles) and 2007 (open circles). The second measurement was made on archived gas splits extracted from the original water samples.

Results and Discussion: Radiocarbon Changes at Shallow and Intermediate Depths

In Figure 2, we compare the 2 cruises (1991 and 2005). In the top 2 panels of Figure 2, we contour the $\Delta^{14}\text{C}$ (in ‰) and in the third panel, the difference (2005–1991) between the gridded fields as $\Delta\Delta^{14}\text{C}$, again in ‰. In the lowest panel, we show a contour map of salinity (in practical salinity units = PSU) for the section. For the bottom 2 panels, we also show potential density anomaly contours (in kg m^{-3}) overlaid in black. The purpose of the latter is to identify the level associated with

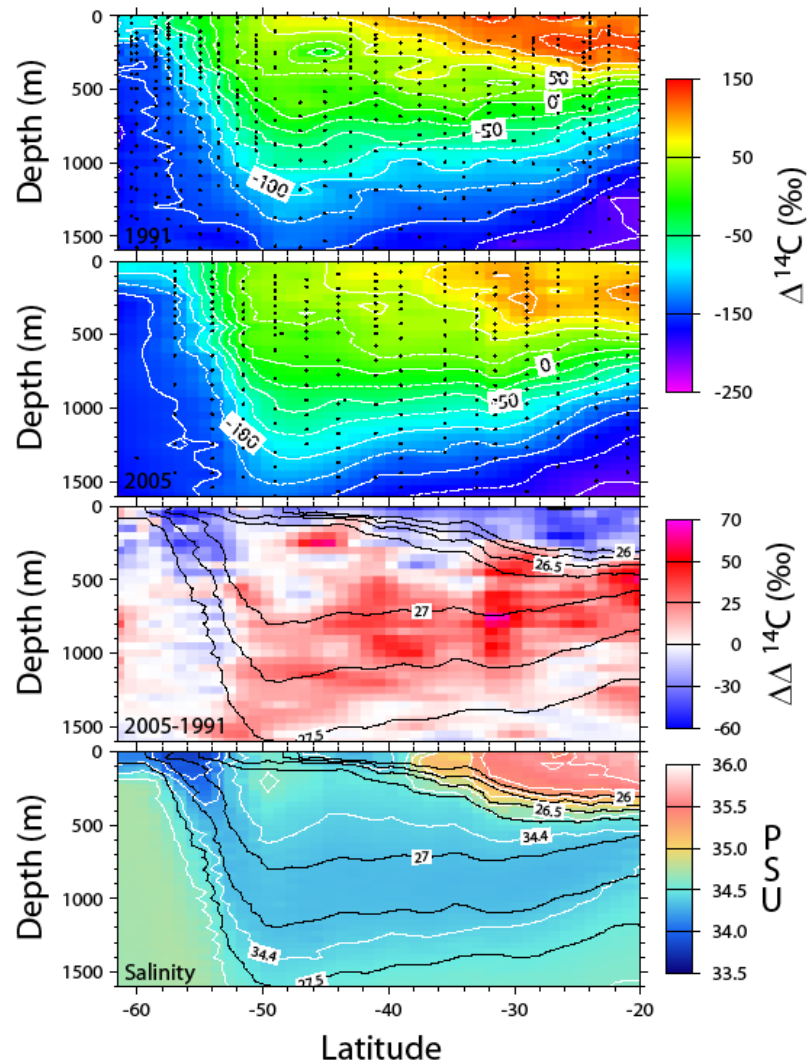


Figure 2 Contour plots (latitude versus depth) of the upper water column distribution of $\Delta^{14}\text{C}$ in 1991 (upper panel) and 2005 (second panel from top), and the difference (2005–1991) between the gridded fields (third panel from top), all in ‰. Sample locations are indicated by black dots in the upper 2 panels. The lowest panel shows the contoured salinity distribution (color map and white contours) with the potential density anomaly (in kg m^{-3}) overlaid as black contours. The potential density anomaly contours are repeated in the ^{14}C difference panel.

the salinity minimum of the Antarctic Intermediate Water that extends downward from the surface at about 56–58°S, deepening and extending northward at depths near ~1000 m at 50°S, shoaling to about 800 m at 20°S.

The 2 ^{14}C sections show similar features, namely a maximum at or near the surface in the subtropics that contrasts with a monotonic decrease in $\Delta^{14}\text{C}$ with increasing depth and latitude. Both sections show a shallow, subsurface ^{14}C maximum equatorward of 30°S: a signature of the STC (Subtropical-Tropical Cell) (McCreary and Lu 1994; Liu and Alexander 2007). Similar features are seen in

meridional sections of both tritium and CFCs, although modified by the space-time character of their introduction into the ocean.

The overall shape of the $\Delta^{14}\text{C}$ distributions (as seen, for example, in the -100‰ contour) mimics the density structure of the subtropical gyre (note the black contours in the lower 2 panels), and reflects the penetration and accumulation of bomb-produced ^{14}C within the thermocline of this oceanographic “province” (Broecker et al. 1985). The $\Delta^{14}\text{C}$ of the deeper waters remains low due to isolation from naturally occurring ^{14}C from the atmosphere as well as the bomb pulse, and in particular because it is ventilated from the Southern Ocean (Kuhlbrodt et al. 2007), which has characteristically high “reservoir ages” (Reimer and Reimer 2001; Butzin et al. 2005) resulting from the generalized upwelling of ancient deep waters along the perimeter of the Circumpolar Current.

Since the ^{14}C stations and sampling depths were not the same for the 2 cruises, we computed the difference field between the 2 cruises by first mapping each section (using Generic Mapping Tool’s SURFACE; Smith and Wessel 1990) onto a $1^\circ \times 50\text{ m}$ rectilinear grid and then subtracting the 2 grids (presented in the third panel of Figure 2). As discussed in the previous section, analytical precision would dictate that differences of $\sim 6\text{‰}$ on individual point comparisons are significant, with averaged differences exceeding 2‰ discernible at the $1\text{-}\sigma$ level. While some of the finer structure in the difference field may in part be an artifact of the gridding process, features larger than the nominal sampling and gridding resolution ($\sim 2^\circ \times \sim 100\text{ m}$) are robust. The color map ranges from -60‰ (blue) to $+70\text{‰}$ (magenta) in $\Delta\Delta^{14}\text{C}$. In the lowest panel of Figure 2, we show the salinity (in PSU, color map and white contours) and for reference we superimpose potential density anomaly contours (in black) on the bottom 2 panels.

Near-surface waters show a systematic decrease in $\Delta^{14}\text{C}$ of approximately $30\text{--}40\text{‰}$ between 1991 and 2005, as would be expected from the observed $\sim 80\text{‰}$ decline in atmospheric $\Delta^{14}\text{C}$ (e.g. see Hua and Barbetti 2004; Q Hua, private communication). The negative surface anomaly penetrates more deeply in the bowl of the subtropical gyre (from approximately 45 to 25°S), and also near the Antarctic Circumpolar Current ($\sim 56^\circ\text{S}$). The former may be rationalized on the basis of Ekman pumping driven subtropical down-welling. The latter is associated with the formation and subduction of Antarctic Mode waters, as evidenced by the penetration of low-salinity waters (e.g. McCartney 1977) depicted in the lowest panel of Figure 2.

The most striking difference, however, is the pronounced increase in $\Delta^{14}\text{C}$ at intermediate depths. The anomaly appears centered on the low-salinity core of the Antarctic Intermediate Water (AAIW), and approaches $50\text{--}60\text{‰}$ at mid-latitudes. We interpret this increase as a downstream manifestation of the evolving ^{14}C anomaly in the Antarctic Intermediate water as it circulates around the South Pacific subtropical gyre. The magnitude of this increase can be used to place rough constraints on both the transit time of freshly ventilated AAIW from the formation region to this section, and the nature of the formation process itself. Coral records show that the predominant bomb- ^{14}C response of the surface ocean was a $\sim 10\text{-yr}$ step-increase that occurred between 1960 and 1970 (Druffel and Suess 1983). Comparison of the observed mid-latitude AAIW increase at this section with the coral responses in a variety of locations strongly suggests that this difference must be due to this transient. Simple one-dimensional advection-diffusion modeling demonstrates that although the magnitude and breadth (over time) of the step-change increase in ^{14}C can be affected by mixing, the advective transport of the centroid of the change is a reflection of the velocity field alone. Given that the observed increase at 150°W apparently occurred sometime between 1991 and 2005, we can estimate the transit time for AAIW of $33 \pm 7\text{ yr}$ from the formation region (i.e. the time difference between 1965 and 1998). Iudicone et al. (2007) identify the formation and entry point of the AAIW in the

southeastern corner of the South Pacific. A crude estimate of the path length along the gyre from this region gives 4000 km. A transit time of 33 yr would require a mean flow rate of approximately $4 \times 10^{-3} \text{ m s}^{-1}$, a number qualitatively consistent with Reid's (1986) geostrophic estimates for the AAIW in this region.

We can also use the 1991 AAIW $\Delta^{14}\text{C}$ observation of -75‰ , corrected for the transit time, to estimate a reservoir age of 600 yr for the source waters of the AAIW. This is somewhat higher than, but not completely inconsistent with, what is obtained from inspection of the results of GCM simulations (Butzin et al. 2005). Inasmuch as AAIW is formed by deep convective events in this region (Georgi 1979; Iudicone et al. 2007), it is likely that the effective reservoir age of this water mass will be increased relative to the regional surface expression.

Results and Discussion: Deep Water Changes

The full depth ^{14}C distributions are presented in Figure 3 (upper 2 panels). They exhibit the expected large-scale ^{14}C characteristics of the abyssal South Pacific, namely a mid-depth core of low- ^{14}C water emanating from the north lying above a tongue of relatively younger (^{14}C -enriched) bottom water entering from the south. Comparison of the full depth ^{14}C differences, contoured in the third panel of Figure 3 shows a surprising large offset in the deep waters. Here again, structural details of the difference field may arise as gridding artifacts, but the overall characteristic of an average increase of 9‰ over the 2000- to 4000-m-depth range between 1991 and 2005 is highly significant relative to possible systematic uncertainties. It should be noted that the offset appears more enhanced near the bottom and less pronounced (perhaps approaching zero) at mid-depths.

The change in ^{14}C appears to be accompanied by changes in hydrography. Although not shown here, the 2005–1991 difference plots of temperature, salinity, dissolved oxygen, and silica along this section in the deep water show a weak but intriguing correlation with the larger-scale structure of the ^{14}C difference plot: higher ^{14}C areas appear associated with more saline, oxygenated, and lower silica waters, and conversely for lower ^{14}C features. Such a correlation might be expected for the intrusion of more recently ventilated waters. The 9‰ overall increase in abyssal ^{14}C suggests this. Calculating differences in the distribution of the Broecker et al. (1999) quantity “phosphate star” ($[\text{PO}_4^*] = [\text{PO}_4] + [\text{O}_2]/175 - 1.95 \text{ } \mu\text{mol kg}^{-1}$) reveals a slight elevation of approximately $0.03 \text{ } \mu\text{mol kg}^{-1}$ in the abyss north of 45°S . An increase in $[\text{PO}_4^*]$ combined with elevated ^{14}C is consistent with an elevated contribution of a “northern source” water (North Atlantic Deep Water) relative to Southern Ocean sources of deep water in the Pacific.

SUMMARY AND CONCLUSIONS

We report a comparison of ^{14}C and hydrographic measurements made along 150°W in the South Pacific in 1991 and 2005. The shallow water ^{14}C patterns show the expected decrease to lower values associated with the atmospheric decline in the bomb ^{14}C transient. The Antarctic Intermediate Waters show a dramatic 50–60‰ increase in $\Delta^{14}\text{C}$ at mid-latitudes, which we attribute to a downstream manifestation of the 1960s oceanic transient response to the atmospheric ^{14}C peak. This allows us to estimate a transit time of order 3 decades between the AAIW formation region and this section, corresponding to advection speeds of order $4 \times 10^{-3} \text{ m s}^{-1}$. The deep waters show a modest but surprising increase in ^{14}C over the intervening years. This increase appears to coincide with small increases in salinity and dissolved oxygen, and a small decrease in dissolved silica. The fact that these are also accompanied by small increase in $[\text{PO}_4^*]$ is suggestive of a shift toward North Atlantic Deep Waters as a source of ventilating waters for the Pacific, consistent with the speculation by Broecker et al. (1999) of possible recent changes in the nature of the global conveyor. The

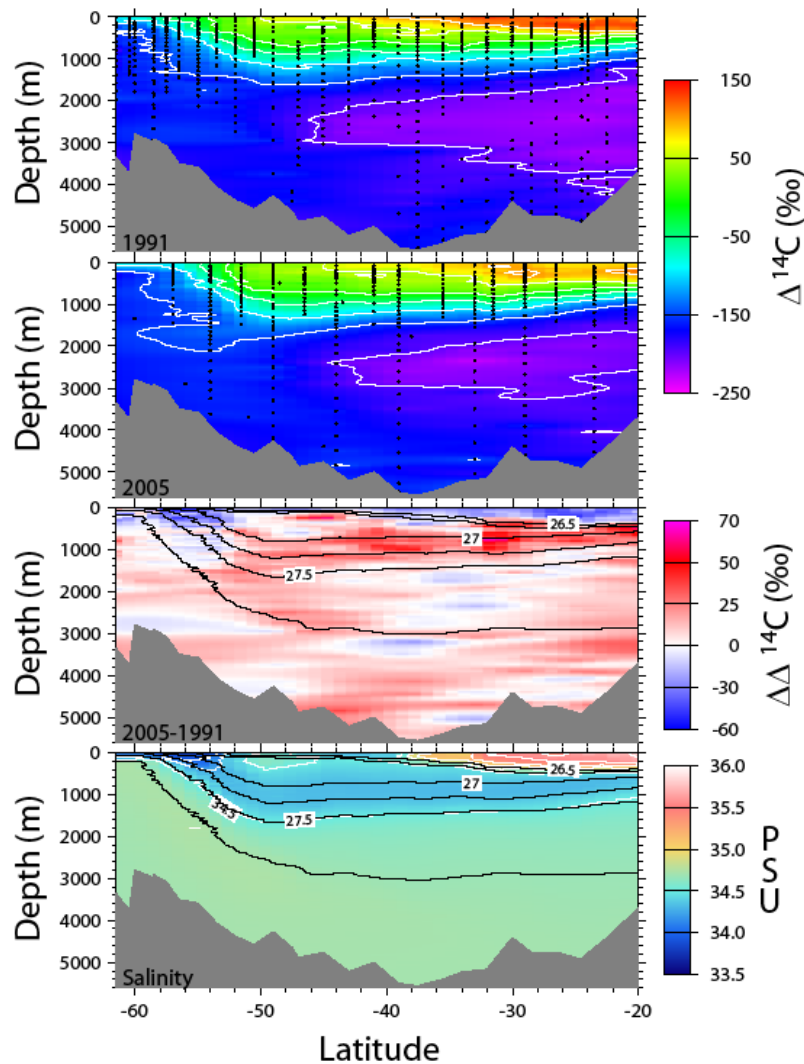


Figure 3 Full depth contour plots (latitude versus depth) of the distribution of $\Delta^{14}\text{C}$ in 1991 (upper panel) and 2005 (second panel from top), and the difference (2005–1991) between the gridded fields (third panel from top), all in ‰. Sample locations are indicated by black dots in the upper 2 panels. The lowest panel shows the contoured salinity distribution (color map and white contours). The potential density anomaly (in kg m^{-3}) is overlaid as black contours on the lower 2 panels.

changes are statistically significant but nonetheless subtle and circumstantial. However, they point to the value both of ^{14}C as a diagnostic of ocean ventilation, and of repeat measurements over the coming decades.

ACKNOWLEDGMENTS

This work was performed under National Science Foundation Grant number OCE-0223434 as well as a cooperative agreement with NSF (most recently OCE-0228996). We are grateful to the Chief Scientists on the WOCE and CLIVAR expeditions, the marine technicians, and the hardworking staff at the National Ocean Sciences AMS Facility.

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